	Application Number		10030232
	Filing Date		2003-08-28
INFORMATION DISCLOSURE	First Named Inventor	Andrew P Bowman	
STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)	Art Unit		2173
	Examiner Name	JOHN	N W. CABECA
	Attorney Docket Numb	er	BO1-0143US

U.S.PATENTS								Remove	
Examiner Initial*	Cite No	Patent Number	Kind Code <sup>1</sup>	Issue E	Date	Name of Patentee or Applicant		Releva	Columns,Lines where nt Passages or Relevant Appear
	1	6467527	B1	2002-1	0-22	Kubota et al.			
	2	6616034	B2	2003-0	9-09	Wu et al.			
	3	6647305	B1	2003-1	1-11	Bigelow			
	4	6749029	B2	2004-0	6-15	Alft et al.			
If you wish	h to a	dd additional U.S. Pater							Add
			U.S.P	ATENT	APPLIC	CATION PUBL	LICATIONS		Remove
Examiner Initial*	Cite No	Publication Number	Kind Code <sup>1</sup>		blication Name of Patentee of Applicant			Releva	Columns,Lines where nt Passages or Relevant Appear
	1	20020188910	A1	2002-1	2-12	Zuzo			
If you wish to add additional U.S. Published Application citation information please click the Add button. Add									
				FOREIG	GN PAT	ENT DOCUM	ENTS		Remove
Examiner Initial*	Cite No	Foreign Document Number <sup>3</sup>	Country Code <sup>2</sup>		Kind Code4	Publication Date	Name of Patente Applicant of cited Document	e or	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear

#### 

	1											
If you wis	h to a	dd add	litional Foreign F	Patent Document	citation	information p	ease o	click the Add	buttor	Add		
				NON-PATE	NT LITE	RATURE DO	CUME	NTS		Remove		
Examiner Initials*	Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposishum, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.						Ţ5					
	1											
If you wish to add additional non-patent literature document citation information please click the Add button Add												
EXAMINER SIGNATURE												
Examiner	Examiner Signature Date Considered											

See Kint Codes of USPTO Patent Documents at Invent/ISPTO.QDV or MPEP 901.04. Enter office that issued the document, by the Nov-letter option Standard ST3.) For Explanates petal or Counters, the Indication of the year of the region or preceded to ever an order to the patent document. A found of countering the Counter of the Emperor may precede to see and not petalent document. If Nord of countering the propriet symbols as indicated on the document under WIPO Standard ST.16 if possible. Sppicant is to place a check mark here if Employin Israquing Personation is attached.

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

# INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10650232		
Filing Date		2003-08-28		
First Named Inventor	Andre	ew P Bowman		
Art Unit		2173		
Examiner Name JOHN		W. CABECA		
Attorney Docket Number		BO1-0143US		

#### CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

	That each item of information contained in the information disclosure statement was first cited in any communication
	from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the
_	information disclosure statement. See 37 CFR 1 97(eV1)

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1/5(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1/7(c)(c)

Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

.7 None

### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/Dale C. Barr, Reg No. 40498/	Date (YYYY-MM-DD)	2006-10-12
Name/Print	Dale C. Barr	Registration Number	40498

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is foll field and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 3T CFR 1.14. This collection is estimated to take it hour to complete, including gathering, preparing and submitting the completed application from to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenan's Office, and Superiment of Commence, P. 0. Bot 1450, Alexandria, V.32.211.450, DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P. 0. Box 1450, Alexandria, V.32.231.4450

## Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
  - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the state of the s
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.